

Amendments To The Claims:

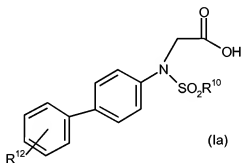
This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

What is claimed is:

1 (Cancelled).

2 (Currently Amended). A compound ~~as claimed in claim 1~~ of formula (Ia):



wherein:

- R¹⁰ represents H or C₁₋₆ alkyl;
 R¹² represents H, halo, CF₃, cyano, OCF₃, nitro, OR¹³, SR¹³, COR¹³ or C₁₋₆ alkyl;
 R¹³ represents C₁₋₆ alkyl or C₁₋₄alkylaryl;
and physiologically functional derivatives thereof.

3 (Cancelled).

4 (Currently Amended). A method for ~~the~~ treatment of a human or animal subject with suffering from or susceptible to an autoimmune disorder or an inflammatory condition, which ~~method~~ comprises administering to said human or animal subject an effective amount of a compound ~~as claimed in of claim 4 2- wherein:~~

the autoimmune disorder or inflammatory condition is selected from:

chronic obstructive pulmonary disease (COPD), asthma, allergen-induced asthmatic reactions, cystic fibrosis, bronchitis, chronic bronchitis, adult respiratory distress syndrome (ARDS), chronic pulmonary inflammation, rhinitis and upper respiratory tract inflammatory disorders (URID), ventilator induced lung injury, silicosis, pulmonary sarcoidosis, idiopathic pulmonary fibrosis, bronchopulmonary dysplasia, arthritis, e.g. rheumatoid arthritis, osteoarthritis, infectious arthritis, psoriatic arthritis, traumatic arthritis, rubella arthritis, Reiter's syndrome, gouty arthritis and prosthetic joint failure, gout, acute synovitis, spondylitis and non-articular inflammatory conditions, herniated/ruptured/prolapsed intervertebral disk syndrome, bursitis, tendonitis, tenosynovitis, fibromyalgic syndrome, inflammatory conditions associated with ligamentous sprain, inflammatory conditions associated with

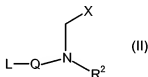
regional musculoskeletal strain, vascular dementia, thrombosis, atherosclerosis, restenosis, reperfusion injury, plaque calcification, myocarditis, aneurysm, stroke, pulmonary hypertension, left ventricular remodeling or heart failure.

5 (Cancelled).

6 (Currently Amended). A pharmaceutical composition comprising a compound **as claimed in of claim 4 2, and a pharmaceutically acceptable carriers or diluents therefor**, and optionally one or more other therapeutic agents selected from anti-inflammatory agents, NSAIDs, beta adrenergic agents, **or** anti-infective agents.

7 (Currently Amended). A process for **the** preparation of compounds of formula (I) **(Ia) as defined in of claim 4 2,** which **process** comprises:

(A) **for the preparation of a compound of formula (I), wherein Z represents a bond and R¹ represents an optionally substituted C₂₋₆alkylaryl or an optionally substituted 5- or 6-membered aryl or heteroaryl,** reacting a compound of formula (II):

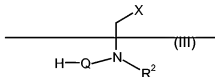


wherein:

R² is SO₂R¹⁰, wherein R¹⁰ is as defined in claim 2;
Q is phenyl; and
X is CO₂H are as previously defined for formula (I); and
L represents a leaving group,

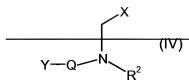
with a reagent suitable to introduce **the group R¹; or phenyl optionally substituted by R¹², wherein R¹² is as defined in claim 2; or**

(B) for the preparation of a compound of formula (I) wherein Z represents a bond and R¹ represents an optionally substituted C₄₋₁₂alkyl, reacting a compound of formula (III):



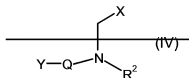
wherein R², Q and X are as previously defined for formula (Ia), with a reagent suitable to introduce the group R¹; or

(C) for the preparation of a compound of formula (I) wherein Z represents O, S, SO, SO₂, NR⁴ or OCR⁶, and R¹ represents an optionally substituted C₄₋₁₂alkyl, reacting a compound of formula (IV):



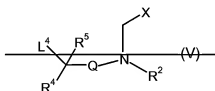
wherein X , R^2 and Q are as previously defined for formula (Ia), and Y represents OH , SH , NR^4H or HCR^4R^5 , or with a reagent suitable to introduce the group R^1 followed in the case where Y is SH by optional oxidation of the sulphide to the sulfoxide or the sulfone; or

(D) for the preparation of a compound of formula (I) wherein Z represents O , S , SO , SO_2 , or NR^4 , and R^1 represents an optionally substituted C_{2-6} alkylaryl or an optionally substituted 5- or 6-membered aryl or heteroaryl, reacting a compound of formula (IV);



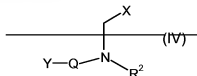
wherein X , R^2 and Q are as previously defined for formula (I), and Y represents OH , SH or NR^4H , with a reagent suitable to couple to the group R^1 , followed in the case where Y is SH by optional oxidation of the sulphide to the sulfoxide or the sulfone; or

(E) for the preparation of a compound of formula (I) wherein Z represents OCR^4R^5 and R^1 represents an optionally substituted C_{2-6} alkylaryl or an optionally substituted 5- or 6-membered aryl or heteroaryl, reacting a compound of formula (V);



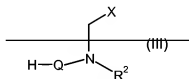
wherein X , R^2 and Q are as previously defined for formula (I) and L^4 is a suitable leaving group, with a reagent suitable to introduce the group R^1-Q ; or

(F) for the preparation of a compound of formula (I) wherein Z represents CR^4R^5O , reacting a compound of formula (IV);



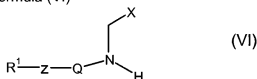
wherein R^2 and Q are as previously defined for formula (I), and Y represents OH, with a reagent suitable to introduce the group $R^1CR^2R^3$; or

(G) for the preparation of a compound of formula (I) wherein Z represents CH_2 , reacting a compound of formula (III):



wherein R^2 , Q and X are as previously defined for formula (I), with a reagent suitable to introduce the group R^1CH_2 ;

(H) reacting a compound of formula (VI)



or a protected derivative thereof, wherein:

R^1 is phenyl optionally substituted by R^{12} ,
wherein R^{12} is as defined in claim 2,
Z is a bond,
Q is phenyl and
X is CO_2H are as previously defined for formula (I),

with a reagent suitable to introduce the group CO_2H R^2 as previously defined for formula (I); or

(J) carrying out a process selected from processes (A) to (G) (H) followed by interconversion of one or more functional groups.

8 (New). The pharmaceutical composition according to claim 6, wherein:

the anti-inflammatory agents are selected from corticosteroids selected from fluticasone propionate, beclomethasone dipropionate, mometasone furoate, triamcinolone acetonide or budesonide);

the NSAIDs are selected from sodium cromoglycate, nedocromil sodium, PDE-4 inhibitors, leukotriene antagonists, CCR-3 antagonists, iNOS inhibitors, tryptase and elastase inhibitors, beta-2 integrin antagonists and adenosine 2a agonists;

the beta adrenergic agents are selected from salmeterol, salbutamol, formoterol, fenoterol or terbutaline and salts thereof; or

the anti-infective agents are selected from antibiotics or antivirals.